

Product Name: Reparixin L-lysine salt

Catalog No.: 6957

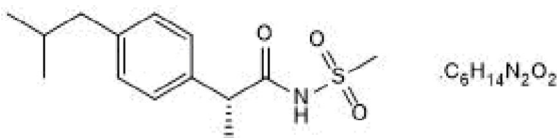
Batch No.: 1

CAS Number: 266359-93-7

IUPAC Name: (αR)- α -Methyl-4-(2-methylpropyl)-*N*-(methylsulfonyl)benzeneacetamide L-Lysine salt

1. PHYSICAL AND CHEMICAL PROPERTIES

| | |
|-----------------------------------|--|
| Batch Molecular Formula: | C ₁₄ H ₂₁ NO ₃ S.C ₆ H ₁₄ N ₂ O ₂ .H ₂ O |
| Batch Molecular Weight: | 447.6 |
| Physical Appearance: | White solid |
| Solubility: | DMSO to 50 mM water to 5 mM with gentle warming |
| Storage: | Store at -20°C |
| Batch Molecular Structure: | |



2. ANALYTICAL DATA

| | |
|---------------------------|--|
| HPLC: | Shows 99.9% purity |
| ¹H NMR: | Consistent with structure |
| Mass Spectrum: | Consistent with structure |
| Optical Rotation: | [α] _D = -21.1 (Concentration = 1.15, Solvent = Methanol) |
| Microanalysis: | |
| | Carbon Hydrogen Nitrogen |
| | Theoretical 53.67 8.33 9.39 |
| | Found 53.41 8.18 9.15 |

Caution - Not Fully Tested • Research Use Only • Not For Human or Veterinary Use

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IUPAC Name: (α R)- α -Methyl-4-(2-methylpropyl)-N-(methylsulfonyl)benzeneacetamide L-Lysine salt

Description:

Reparixin L-lysine salt is a potent and noncompetitive CXCR1 and CXCR2 allosteric antagonist (IC_{50} = 1 nM for inhibition of CXCL8-induced human polymorphonuclear cell migration). Also inhibits migration of rodent neutrophils induced by CXCL1, CXCL2, CXCL8 and CINC-1. Inhibits vascular permeability and neutrophil recruitment in in vivo models of mild and severe ischemia/reperfusion injury. Also selectively depletes cancer stem cells in human breast cancer cell lines and xenograft models.

Physical and Chemical Properties:

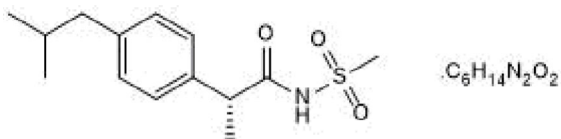
Batch Molecular Formula: $C_{14}H_{21}NO_3 \cdot C_6H_{14}N_2O_2 \cdot H_2O$

Batch Molecular Weight: 447.6

Physical Appearance: White solid

Minimum Purity: $\geq 98\%$

Batch Molecular Structure:



Storage: Store at $-20^{\circ}C$

Solubility & Usage Info:

DMSO to 50 mM

water to 5 mM with gentle warming

This product is supplied in lyophilized form. It may appear as a solid, gel or film and be very hard to visualize. Solutions should be made by adding solvent directly to the vial. The vial should then be vortexed vigorously to ensure the product has completely dissolved.

Stability and Solubility Advice:

Some solutions can be difficult to obtain and can be encouraged by rapid stirring, sonication or gentle warming (in a $45-60^{\circ}C$ water bath).

Information concerning product stability, particularly in solution, has rarely been reported and in most cases we can only offer a general guide. *Unless contradicted by product-specific protocols or instructions, our standard recommendations apply:

SOLIDS: Provided storage is as stated on the product label and the vial is kept tightly sealed, the product can be stored for up to 6 months from date of receipt.

SOLUTIONS: We recommend that stock solutions, once prepared, are stored aliquoted in tightly sealed vials at $-20^{\circ}C$ or below and used within 1 month. Wherever possible solutions should be made up and used on the same day.

References:

Ginestier et al (2010) CXCR1 blockade selectively targets human breast cancer stem cells *in vitro* and in xenografts. *J.Clin.Invest.* **120** 485. PMID: 20051626.

Bertini et al (2004) Noncompetitive allosteric inhibitors of the inflammatory chemokine receptors CXCR1 and CXCR2: prevention of reperfusion injury. *Proc.Natl.Acad.Sci.U.S.A.* **101** 11791. PMID: 15282370.

Souza et al (2004) Repertaxin, a novel inhibitor of rat CXCR2 function, inhibits inflammatory responses that follow intestinal ischaemia and reperfusion injury. *Br.J.Pharmacol.* **143** 132. PMID: 15302676.

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